Amendments to the Claims

This Listing of Claims will replace all prior versions and listings of claims in the application.

Listing of Claims

1. -2. (Cancelled)

- 3. (Currently Amended) A vaccine composition comprising a proteosome adjuvant and at least onea group A Streptococcal antigen attached to a hydrophobic moiety, wherein the antigenthat comprises an antigenic peptide between 6 and 25-15 and 30 amino acids in length from the conserved C-terminal region of an S. pyogenes M protein, wherein the antigenic peptide comprises the amino acid sequence ASREAKKQVEKALE (SEQ ID NO:1), and wherein the antigen is attached to a hydrophobic moiety.
- 4. (Currently Amended) The vaccine composition according to claim 3 wherein the antigenic peptide has consists of the sequence ASREAKKQVEKALE (SEQ ID NO:1).
- 5. (Currently Amended) The vaccine composition according to claim 3 wherein the <u>antigen comprises the antigenic peptide is</u>—flanked by amino acid sequences to maintain helical folding of the antigen.
- 6. (Currently Amended) The vaccine composition according to claim 5 wherein the <u>antigenic peptide has antigen comprises</u> the sequence KQAEDKVKASREAKKQVEKALEQLEDKVK(SEQ ID NO:2).

7. (Cancelled)

- 8. (Previously Presented) The vaccine composition according to claim 3 wherein the hydrophobic moiety is attached at the N-terminal end or C-terminal end of the antigen for complexing the antigen with the proteosome adjuvant.
- 9. (Currently Amended) The vaccine composition according to either-claim 3 or claim 7 for wherein the composition is formulated for mucosal administration.
- 10. (Previously Presented) The vaccine composition according to claim 9 wherein mucosal administration is intranasal.

11. - 12. (Cancelled)

- 13. (Previously Presented) The vaccine composition according to claim 3 wherein administration of the composition to an individual induces a mucosal immune response.
- 14. (Previously Presented) The vaccine composition according to claim 3, wherein administration of the vaccine composition induces a serum immune response.
- 15. (Previously Presented) The vaccine composition according to claim 9 wherein the vaccine composition is capable of treating or preventing a group A Streptococcal infection via reducing or preventing streptococcal group A bacterial colonisation of the throat.
- 16. (Currently Amended) A method of treatment or prophylaxis of group A Streptococcal infection in an individual comprising administering the vaccine composition according to either-claim 3 or claim 7 to the individual.
- 17. (Previously Presented) The method according to claim 16 wherein said vaccine composition is administered intranasally to said individual.

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- 18. (Previously Presented) The method according to claim 17 wherein the treatment or prophylaxis of the group A Streptococcal infection is produced via prevention or reduction of bacterial colonisation of the throat.
- 19. (Previously Presented) The vaccine composition according to claim 3 wherein the antigen further comprises a spacer peptide comprising at least two glycine residues, and wherein the spacer peptide links the antigenic peptide and the hydrophobic moiety.